

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

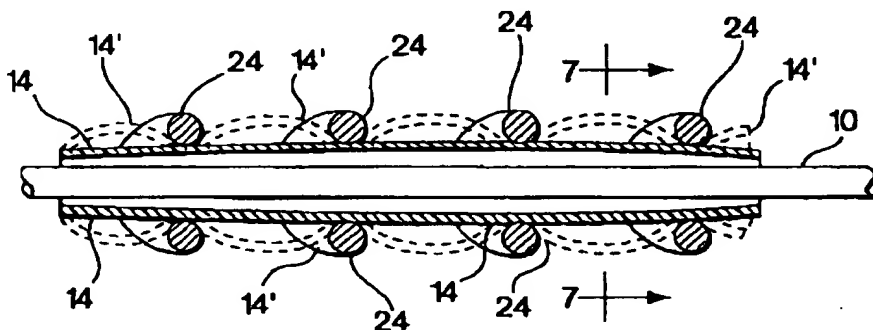
**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problems Mailbox.**



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶: A61F 2/06	A1	(11) International Publication Number: WO 99/15106 (43) International Publication Date: 1 April 1999 (01.04.99)
(21) International Application Number: PCT/IE98/00076 (22) International Filing Date: 16 September 1998 (16.09.98) (30) Priority Data: 08/934,703 22 September 1997 (22.09.97) US (71) Applicant: BARD CONNAUGHT [IE/IE]; 2 Harbourmaster Place, Custom House Dock, Dublin 1 (IE). (72) Inventors: WHELAN, Sean; Cloonacauneen, Castlegar, County Galway (IE). PURCELL, Neil; 58 Churchfields, Lower Salthill, Galway (IE). FITZMAURICE, Thomas; Oranswell, Bushy Park, Galway (IE). (74) Agent: MACLACHLAN & DONALDSON; 47 Merrion Square, Dublin 2 (IE).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>

(54) Title: BALLOON MOUNTED STENT AND METHOD THEREFOR



(57) Abstract

The present invention concerns a mechanism for securely mounting an expandable stent onto a balloon catheter for intraluminal delivery within a patient. The stent is secured to the balloon of the delivery catheter by forming the surface of the balloon to project into the interstices of the stent. To mount the stent it is first slid over an evacuated and wrapped balloon while in its compact delivery diameter. A rigid tube then is placed over the stent and balloon assembly and the balloon is pressurized while heat is applied. The rigid tube prevents the stent from expanding but allows the balloon to deform so that its surface projects slightly through either or both of the interstices and ends of the stent. The elevated temperature causes the balloon material to take a permanent set into the stent such that once pressure is removed, the stent is interlocked with the surface of the balloon. The engagement of the stent with the balloon prevents unintended axial sliding of the stent with respect to the balloon during delivery through the vessel of the patient.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NI	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

- 1 -

BALLOON MOUNTED STENT AND METHOD THEREFOR

Field of the Invention

This invention relates to stents for placement in the human body and delivery systems therefor.

Background of the Invention

5 The use of intraluminal stents has had increasing acceptance and use in the treatment of various medical conditions. Intraluminal stents can be used to maintain the patency of blood vessels and other body lumens, such as in the treatment of urological disorders, among others. The use of stents to maintain open a coronary
10 or other artery after an angioplasty procedure has been performed has become a common practice. Over the past decade, various stent designs have been described and used. Many have been of the type in which the stent is in a tubular configuration that can be expanded from a relatively small diameter adapted for delivery through the vasculature (low profile) to a larger deployed diameter by
15 mounting the stent on the balloon of an inflatable balloon catheter and then advancing the catheter to place the balloon mounted stent at the intended site of deployment. The balloon then is inflated to expand the stent into engagement with the body lumen. The stent maintains its expanded shape to support the body lumen. The balloon then is deflated and the delivery catheter is withdrawn, leaving
20 the stent in place to support the body lumen. The configuration of such stents typically may be considered as generally tubular in which the wall of the tube is defined by variously configured wires or a tubular member that has been shaped to provide a plurality of interconnected wire-like struts or wires. Illustrative examples of such stents may be found in U.S. patents nos. 4,733,665 (Palmaz), 5,104,404
25 (Wolff), 5,421,955 (Lau) and in International Publication WO 96/41591 (Borghi), among others.

The design and placement of a stent can present a number of considerations, particularly when the stent is to be used in an environment where it must be

- 2 -

advanced through tortuous anatomy, as is not uncommon in coronary angioplasty. The stent must have the ability to maintain its tubular configuration when expanded in the body lumen in order to maintain the body lumen open. It also should have sufficient longitudinal flexibility, when mounted in its low profile on the balloon, so
5 that it can be advanced through curved, sometimes tortuous, blood vessels in order to reach the deployment site. Additionally, the position of the stent on the balloon should be maintained without shifting longitudinally, as the balloon mounted stent is advanced to the deployment site. That can present particular difficulty when the stent and balloon to which it is mounted must be passed through sharply curved
10 tortuous anatomy or anatomy where the lumen has become narrowed or otherwise partly obstructed regions. Should the stent engage a blood vessel in a sharply curved or somewhat narrowed region and consequently, shift longitudinally on the balloon as the catheter is advanced or manipulated, the balloon, when inflated, may not expand the stent fully along the length of the stent. In some cases, the stent
15 could be dislodged from the balloon without the ability to retrieve it.

The importance of maintaining the stent securely on the balloon has been recognized in the prior art. A number of approaches have been proposed. Some simply slide the stent onto the balloon and crimp it tightly about the balloon. That may risk damage to the balloon or the stent and also may result in a relatively
20 longitudinally stiff stent. Other stent retention devices have been proposed including the use of a sleeve to overlie the stent during advancement, with the sleeve being retracted when the stent is to be deployed. Still other approaches have included the use of end caps mounted on the catheter and adapted to temporarily engage the ends of the stent while permitting the stent ends to release when the stent is
25 expanded as is shown in U.S. patent no. 4,950,227 (Savin). Such devices and techniques generally have required compromises of one or more desirable features of the stent. The desirable low profile or longitudinal flexibility may be compromised as well as the security of retention. It would be desirable to provide a balloon mounted stent and method for such mounting that would reduce the extent of
30 compromise of desirable characteristics of the stent and its delivery system.

- 3 -

Summary of the Invention

In accordance with the invention, a stent and balloon catheter assembly is provided in which after the stent is mounted on the balloon that has been wrapped about the catheter shaft in a relatively tight, low profile, the balloon is molded against
5 the stent to secure the stent directly to the balloon. The outer diameter of the stent is maintained during the process, as by placing a rigid retaining tube about the stent. The interior of the balloon then is pressurized sufficiently so that, although wrapped in a low profile configuration, it will tend to expand radially outwardly against the stent. Concurrently with such pressurization, polymeric material of the balloon is
10 softened, as by heating, exposure to a solvent or other means for softening the balloon sufficiently to enable it to be molded. The balloon material may be softened by any appropriate means including heat or solvents. The soft moldable condition of the balloon is maintained sufficiently to enable portions of the balloon to expand and mold against the ends of the stent or into interstices along the length of the stent, or
15 both. Portions of the balloon thus are expanded to a radius that is greater than the radius of the inner lumen defined by the stent. The balloon is maintained under conditions of pressure and softened condition for a predetermined time to assure the balloon will maintain its molded state. In the resulting assembly, the stent is interengaged with portions of the balloon to provide substantial resistance to shifting
20 of the longitudinal position of the stent along the balloon. The resistance to longitudinal shifting created by the interengaged portions supplements frictional engagement between the stent and balloon serving to resist their relative axial movement. The balloon, being longer than the stent may expand beyond the ends of the stent toward the inner diameter of the retaining tube to form a shoulder that
25 engage the ends of the stent to provide resistance to axial movement. The balloon expansion at the ends of the stent also reduces the abruptness of the transition between the balloon and the end of the stent to reduce the risk of snagging the stent on the guide catheter or vessel.

A general object of the invention is to provide an improved system for
30 mounting a stent securely onto the balloon of a stent delivery catheter without

- 4 -

increasing the profile of the balloon and stent combination. Also among the objects of the invention are to provide an improved balloon-stent combination in which the stent is secured on the balloon of a stent delivery catheter by interfering engagement of the stent directly with the balloon; to provide a balloon catheter and
5 mounted stent adapted to pass through tortuous anatomy with reduced risk of the stent being displaced from its initial position on the balloon; to provide an improved technique for mounting a stent on the balloon of a stent delivery catheter; to provide a technique for effecting direct engagement between a stent and the balloon of a
10 delivery catheter in which the stent is securely mounted on the balloon without use of supplemental retaining or restraining devices; to provide an assembly of a stent mounted securely on the balloon of a balloon catheter without significantly affecting the longitudinal flexibility of the assembly; and to provide a method for mounting a stent on the balloon of a stent delivery catheter in which some portions of the balloon project radially beyond the radius defined by the inner surface of the stent.

15

Description of the Drawings

The foregoing and other objects and advantages of the invention will be appreciated more fully from the following further description thereof with reference to the accompanying drawings wherein:

20

FIG. 1 is a diagrammatic illustration of a stent delivery catheter having a stent mounted on the balloon of a delivery catheter;

FIG. 1A is cross-sectional view of the catheter shown in FIG. 1 as seen along the line 1A-1A;

FIG. 1B is a diagrammatic illustration of the distal end of the catheter showing the configuration of the balloon, when inflated, and the shaft of the catheter;

25

FIG. 2A is a diagrammatic transverse cross-sectional illustration of the balloon region of a balloon catheter with the balloon having been evacuated to form radially extended wings;

FIG. 2B is a diagrammatic illustration similar to FIG. 2A in which the wings of the evacuated balloon have been wrapped about the catheter shaft to a low profile

- 5 -

configuration;

FIG. 2C is a diagrammatic illustration similar to that of FIGS. 2A and 2B but in which the balloon has been inflated;

FIG. 3 is a diagrammatic illustration of a wrapped balloon and a stent
5 mounted on a loading tube in readiness to be placed on the balloon;

FIG. 4 is an illustration of the balloon region of the stent delivery catheter with the stent having been located on the balloon and with a tubular restraining member being advanced over the assembly of the stent and balloon;

FIG. 5 is a diagrammatic, longitudinal sectional illustration of the interfacing
10 region between the balloon and the stent as seen along the line 5-5 of FIG. 1 in which the configuration of the interfacing region before pressure molding is shown in solid lines, and the configuration of the interfacing region after pressure molding is shown in phantom;

FIG. 5A is a diagrammatic longitudinal sectional illustration of the manner in
15 which the balloon can be molded against the ends of a stent to form shoulders engageable with the ends adapted to resist longitudinal movement of the stent on the balloon;

FIG. 6 is a diagrammatic transverse section through the stent illustrating the relationship of the inner radius, outer radius and virtual wall thickness defined by the
20 stent; and

FIG. 7 is a transverse sectional illustration of the balloon mounted stent as seen along the line 7-7 of FIG. 5 showing the manner in which portions of the balloon protrude radially beyond the inner radius of the stent and into engagement with radially extending surfaces of the stent.

25 Description of the Illustrative Embodiment

The invention, for convenience, is described in the context of a stent adapted for delivery in conjunction with an angioplasty procedure, although it should be understood that the invention may be practiced in connection with stents and catheters adapted for other medical procedures. In the preferred embodiment, the

- 6 -

stent delivery catheter includes a catheter shaft 10 having at least one lumen 12 for inflation of a balloon 14 mounted at the distal region of the catheter shaft 10. As shown in FIG. 1B, the balloon may be configured to have a generally cylindrical, uniform diameter central section 11 and a pair of outwardly tapering end cones 13, each of which terminates in a neck 15 by which the balloon is attached to the catheter shaft 10. The portion of the catheter shaft 10 that extends through the balloon may be provided with a pair of longitudinally spaced radiopaque markers 17. The markers may be disposed on the shaft 10 at approximately the regions where the end cones 13 merge into the central balloon section 11.

The catheter shaft also may have a guidewire lumen 16 to enable the catheter to be advanced and guided to the deployment site along a guidewire 18. The proximal end of the shaft 10 is provided with a fitting 20 that includes a connector 22 by which the inflation lumen 12 can be connected to a balloon inflation device (not shown). While a dilatation catheter intended for angioplasty may be used in the practice of the present invention, it is not necessary that the balloon 14 be suitable for an angioplasty procedure as it need only be capable of retaining and expanding the stent in a delivery procedure that is separate from the angioplasty procedure. However, the balloon of the stent delivery catheter of the present invention should have characteristics that enable it to be molded into engagement with the interstices or other radially extending surfaces of the stent. For example, in the illustrative embodiment, the balloon may be that described in U.S. patent 5,500,180 (Anderson et al.).

In accordance with the invention a stent 24 having a generally tubular configuration is mounted on and about a balloon 14. The stent may be somewhat shorter than the cylindrical central section 11 of the balloon to assure that when the balloon is inflated within the patient to deploy the stent, the stent will be expanded along its length by the balloon.

The invention may be practiced with any balloon-expandable stent. For purposes of illustration, the invention is described as being used in connection with a stent described in International Patent Application, Publication WO 96/41591

- 7 -

(Borghi), the disclosure of which is hereby incorporated by reference, in its entirety. Other examples of balloon expandable stents include those disclosed in U.S. patent Nos. 4,733,665 (Palmaz), 5,041,126 (Gianturco) and 5,104,404 (Wolff), the disclosures of which also are hereby incorporated by reference, in their entireties. In the illustrative embodiment, the stent is characterized by an arrangement of wire-like hoops 23 connected to a spine 25 that, taken together, may be considered to define a virtual cylindrical tubular shape. The reticulated wires or struts 26 that form the stent are arranged to define spaces 27 along the length of the stent, the configuration and orientation of which may vary as a function of the geometry and design of the stent. The tubular stent thus may be considered as defining an inner cylindrical radius R_i and an outer cylindrical radius R_o , the difference between the two radii defining what may be considered as the wall thickness T of the stent (FIG. 6). The stent may be formed by a number of techniques familiar to those skilled in the art and may be formed from a variety of materials, including metals such as stainless steel or nickel titanium alloys (e.g., nitinol), polymeric or bioabsorbable materials. The structure and material of the stent should be capable of enabling expansion from its low profile configuration, in which it is adapted to be advanced through the vasculature to the deployment site, to an expanded diameter in which it can engage and support the wall of a blood vessel or other body lumen, as appropriate.

The stent is intended to be mounted on a balloon that itself has been configured in a low profile. As shown in FIGS. 2A-2C, a common technique for configuring the balloon in a low profile is first to evacuate the balloon to cause the balloon to assume a cross-sectional shape as shown in FIG. 2A in which the balloon is collapsed and defines a plurality of radially extending flat wings 28. While maintaining the evacuation pressure, the wings of the balloon are wrapped closely about the catheter shaft in one of several compact configurations to maintain a low profile about the shaft, such as an S-shaped configuration shown in FIG. 2B.

With the balloon in its wrapped, low profile configuration, the stent then is loaded onto the wrapped balloon as suggested in FIG. 3. The stent may be

- 8 -

positioned on the balloon so that the ends of the stent are disposed adjacent and between the radiopaque marker bands on that portion of the catheter shaft that passes through the balloon. To facilitate loading, the stent may be mounted about and crimped onto a thin wall polymeric tube, shown in phantom at 29 that, in turn, can be slipped over the balloon. The tube 29 is advanced to position the stent in the desired location relative to the balloon with the end cones 13 protruding beyond the ends of the stent. Then, while holding the stent in the desired position relative to the balloon, the tube 29 is withdrawn, leaving the stent in the desired longitudinal position about the balloon. The stent may be crimped slightly by hand or with a crimping tool to achieve a snug fit about the balloon.

A relatively rigid tube 30 then is placed over the stent. The tube 30 may be formed from a suitably rigid tubing such as a polymer commercially available under the trade designation KYNAR and is intended to restrain the stent from radial expansion during the remainder of the stent mounting process. The tube 30 should be of sufficient length to completely cover the stent and the balloon and may be of the order of twice as long as the stent. The tube 30 should be dimensioned such that a relatively small clearance exists between the inside diameter of the tube and the outside surface of the stent as, for example, on the order of 0.001" - 0.003" to facilitate placement of the tube 30 over the stent. The balloon then is pressurized by connection of a suitable device to the balloon inflation leg of the proximal fitting. In the illustrative example, the balloon may be pressurized with a gas, such as air or nitrogen, within the range of 6 to about 20 bar, preferably about 10 bar. Although the balloon has been wrapped tightly in a low profile configuration, the wrapping cannot completely seal off all minute flow passages through the balloon. Consequently, it is possible to apply sufficient pressure within the wrapped balloon to cause portions of the balloon to press radially outwardly against the stent.

In the illustrative embodiment, the assembly of the pressurized balloon catheter, stent and molding tube 30 then is heated to raise the temperature of the balloon material sufficiently to allow it to yield and be molded around the ends and against the interior of the stent. A temperature in the range of 50° - 70° C, preferably

- 9 -

65°C, under the above pressures has been found to be sufficient to cause the balloon to become molded to the stent. Although the expansion of the balloon may be slight, it is sufficient to cause some portions of the balloon to expand to a radius that is greater than the inner radius R_i of the stent so that those portions of the balloon may be considered as projecting radially outwardly into the region of the virtual wall thickness T . The conditions of temperature and pressure are maintained sufficiently long to assure that the balloon will set and remain in that configuration when the balloon is cooled and pressure is removed. FIG. 7 is a transverse sectional illustration of the balloon mounted stent after the balloon has been molded and expanded radially outwardly so that portions of the balloon, indicated at 14' protrude radially outwardly beyond the inner radius R_i of the stent as defined by the struts 26. FIG. 5 is a longitudinal illustration of the balloon and stent of FIG. 7. FIG. 5A illustrates, diagrammatically and in enlarged scale, the region of the molded balloon with the end of a stent as confined within the retaining tube. The shoulder thus formed enables the balloon to engage directly the ends of the stent to resist longitudinal movement of the stent on the balloon. This embodiment may be used with any type of stent, including stents that have few or no interstices when in the low profile configuration.

Applicants have found that, in the illustrative embodiment, applying heat and temperature for a time of approximately 30 - 90 seconds, preferably about 60 seconds, sufficiently molds the balloon to the stent. A brief cooling period of approximately one minute immediately following the heating period, while maintaining pressure at 10 bar also may help set the balloon in its molded configuration. Although the balloon and stent may expand slightly to contact the inside of the tube 30 while pressure is applied, slight elastic recoil of the stent after the inflation pressure is removed results in a slight clearance on the order of 0.0005" between the tube and the stent. The restraining tube 30 then is easily removed and the stent and balloon may be recompressed about the catheter shaft by hand, or with a suitable tool to reduce the slight expansion that occurred during the balloon/stent engagement process. The delivery catheter with premounted stent

- 10 -

then may be packaged in readiness for use in a procedure.

It should be understood that although, in the illustrative embodiment, the balloon material is softened by the application of heat, other procedures may be employed to soften the balloon in the practice of the invention. For example, the
5 balloon material may be softened, where appropriate, with a solvent sufficiently and under conditions to enable the balloon to be molded under the influence of the pressure differential from the interior to the exterior of the balloon.

From the foregoing, it will be appreciated that the invention provides a balloon mounted stent and a method for mounting the stent on the balloon in which there are
10 no stent retaining components to increase the profile of the assembly while maintaining the stent in a secure position on the balloon by direct, interfering engagement of the stent with the balloon. The invention provides a balloon catheter and mounted stent in which the longitudinal flexibility is not adversely affected and that can pass through tortuous or restricted anatomy with reduced risk of the stent
15 becoming dislodged or displaced from its position on the balloon. A number of advantages thus are achieved with minimal compromise. It should be understood, however, that the foregoing description of the invention is intended merely to be illustrative thereof and that other embodiments, modifications and equivalents within the scope of the invention may be made by those skilled in the art.

20 Having thus described the invention, what we desire to claim and secure by Letters Patent is:

- 11 -

Claims

1. A stent and delivery catheter therefor comprising, in combination:
a catheter having a shaft and an inflatable balloon at its distal end;
a stent disposed on and about the balloon, the stent defining an inner
5 diameter;
at least part of the balloon projecting radially beyond the inner diameter
of the stent to resist longitudinal movement of the stent on the balloon.
2. A stent and catheter combination as defined in claim 1 further
comprising:
10 the stent having a plurality of spaces defined between its ends, the
radially projecting portions of the balloon projecting into at least some of the spaces.
3. A stent and catheter combination as defined in claim 1 further
comprising:
the balloon being wrapped circumferentially about the catheter shaft.
- 15 4. A stent and catheter combination as defined in any one of claims 1-3
wherein the part of the balloon that projects radially beyond the inner diameter of the
stent comprises a portion of the balloon disposed immediately adjacent at least one
end of the stent and formed to define a shoulder engageable with the end of the
stent.
- 20 5. A stent and delivery catheter therefor comprising, in combination:
a catheter having an inflatable balloon;
a tubular stent mounted on the balloon;
the balloon having portions molded to engage the stent to resist
relative longitudinal sliding movement of the stent with respect to the balloon.

- 12 -

6. The combination as defined in either one of claims 1 or 5 wherein the region of the assembly of the catheter, balloon and stent have substantially the same longitudinal flexibility as that which the assembly would have in the absence of the balloon-to-stent engagement for resisting relative longitudinal movement.

5 7. A stent as defined in either one of claims 1 or 5 wherein the engagement of the balloon with the stent comprises the sole means of attaching the stent to the balloon to resist relative longitudinal movement.

8. A stent and delivery catheter therefor comprising, in combination:
a catheter having a shaft and an inflatable balloon at its distal end;
10 a stent on and about the balloon;
means for directly interengaging the stent with the balloon to resist longitudinal movement of the stent relative to the balloon.

9. A stent and delivery catheter therefor comprising, in combination:
a catheter having a shaft and radially expandable member at its distal
15 region;
a tubular stent mounted on the expandable member;
at least part of the expandable member and the stent being
interengaged sufficiently to resist longitudinal movement of the stent on the
expandable member, said interengagement comprising the primary means for
20 resisting said longitudinal movement.

10. A method for mounting a stent on a balloon of a catheter comprising:
configuring the balloon in a low profile;
placing a balloon expandable stent on the balloon;
urging portions of the balloon radially outward to a radius greater than
25 the inner radius of the stent to resist longitudinal movement of the balloon and stent
while in their low profile.

- 13 -

11. A method as defined in claim 10 wherein the step of urging the balloon radially outwardly comprises molding at least part of the balloon radially outwardly into engagement with and against the inner contour of the stent.

5 12. A method as defined in claim 11 wherein the molding step further comprises:

after placing the stent on the balloon, concurrently applying pressure to the interior of the balloon and softening the balloon material sufficiently to enable it to be molded against the inner contour of the stent under the influence of said pressure;

10 maintaining the softened balloon in the pressurized condition, sufficiently to cause portions of the balloon to be molded radially outwardly against and in conformity to at least a portion of the inner contour of the stent.

13. A method as defined in claim 11 wherein said step of softening the balloon material comprises applying heat to the balloon material.

15 14. A method as defined in claim 12 wherein the step of softening the balloon material comprises applying a solvent to the balloon material.

15. A method as defined in claim 12 wherein the step of softening the balloon material comprises exposing the balloon to a means for softening the balloon material.

20 16. A method as defined in any one of claims 12-15 further comprising:
cooling the balloon while maintaining the pressure in the balloon,
allowing the balloon to set to a molded configuration corresponding in part to the inner contours of the stent.

- 14 -

17. A method as defined in any one of claims 12-15 further comprising:
while maintaining the pressure in the balloon, allowing the balloon to set to a molded configuration that includes a shoulder adjacent and engageable with at least one end of the balloon.

5 18. A method as defined in claim 12 further comprising maintaining the diameter of the stent while the balloon is pressurized.

19. A method as defined in claim 18 wherein the step of maintaining the stent diameter comprises placing a restraining member about the exterior of the stent before the interior of the balloon is pressurized.

10 20. A method as defined in claim 19 wherein the step of maintaining the diameter of the stent comprises placing a rigid tubular member over the stent.

1/3

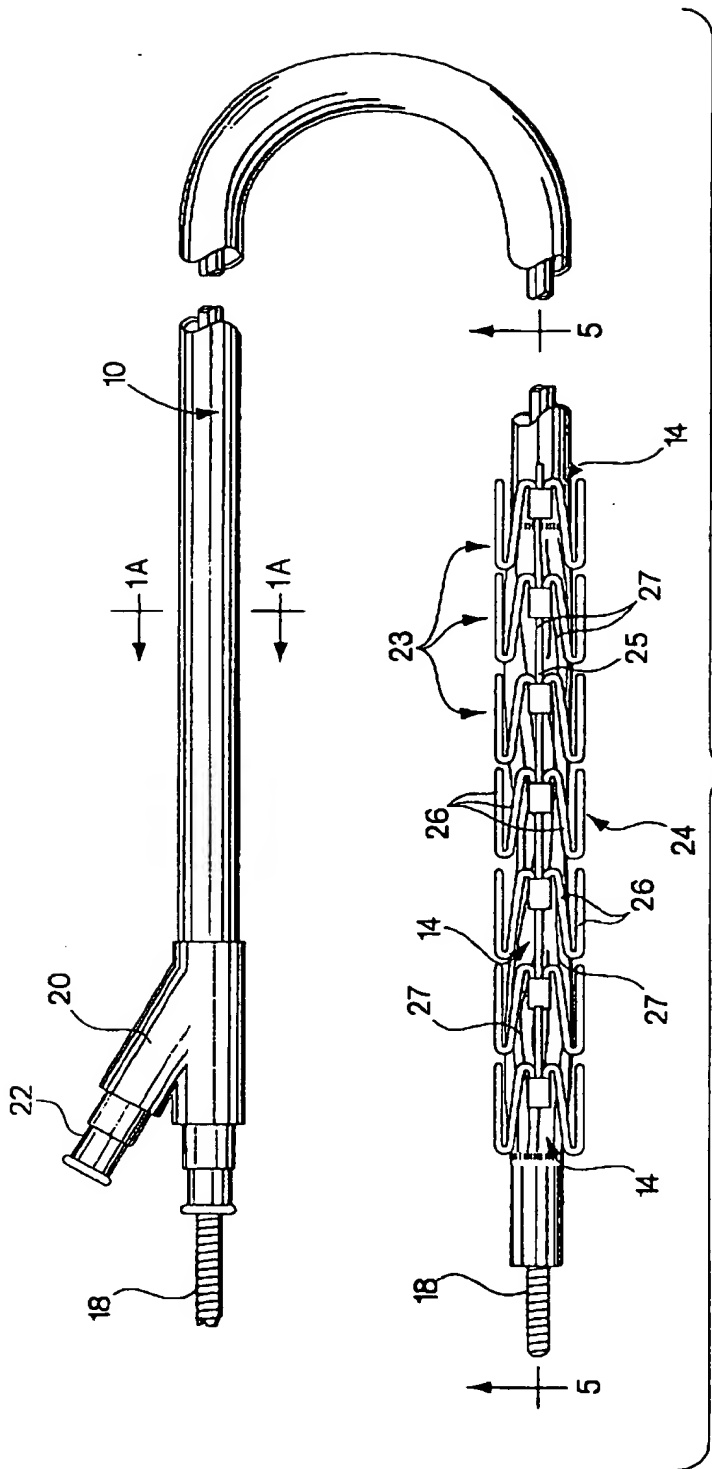


Fig. 1

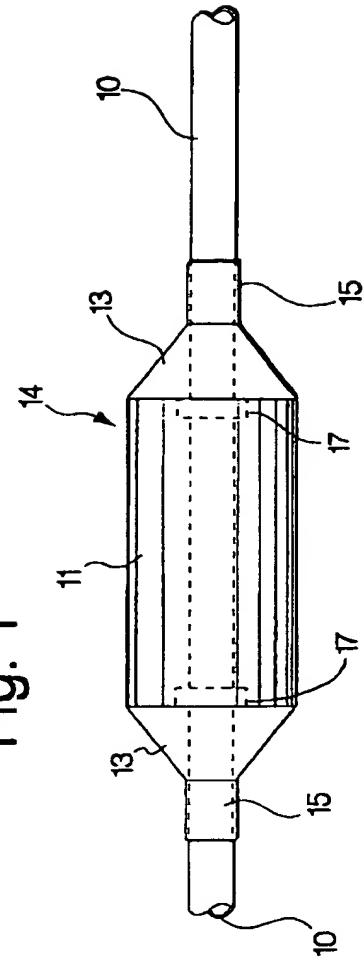


Fig. 1A

Fig. 1B

SUBSTITUTE SHEET (RULE 26)

2/3

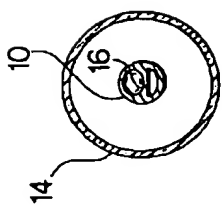


Fig. 2C

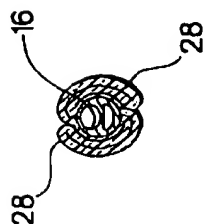


Fig. 2B

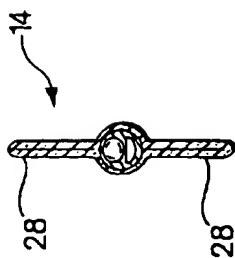


Fig. 2A

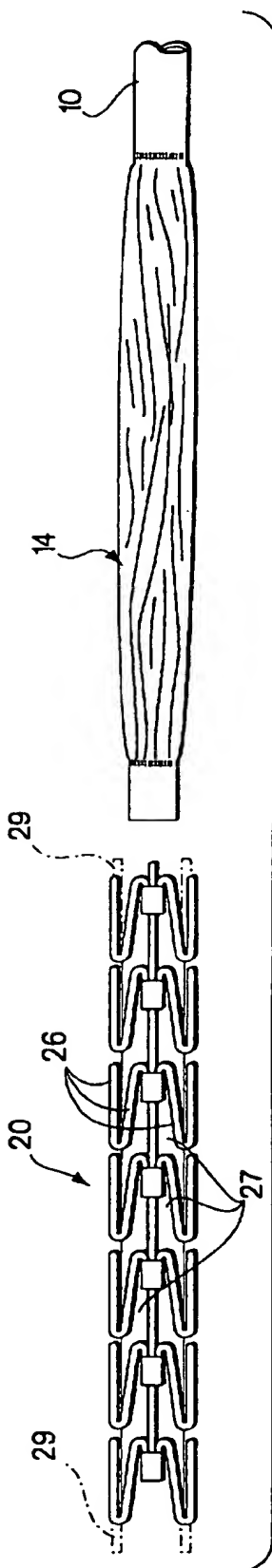


Fig. 3

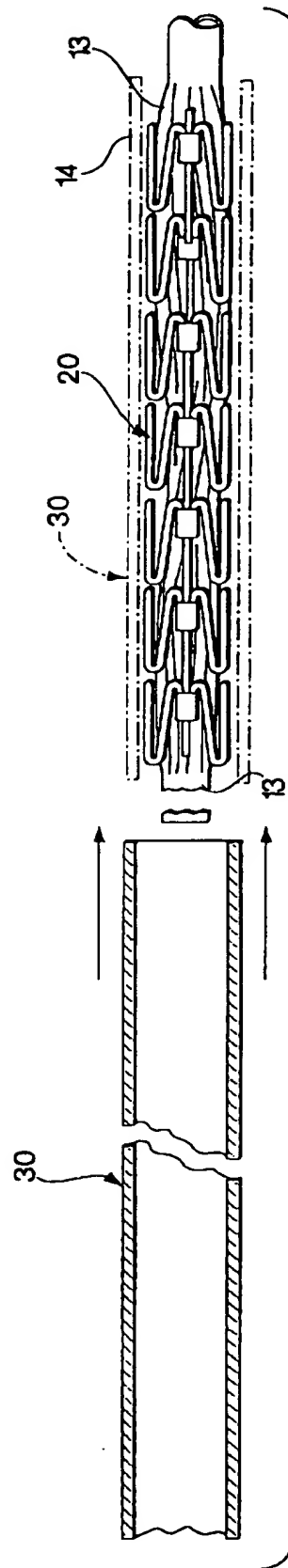


Fig. 4

SUBSTITUTE SHEET (RULE 26)

3/3

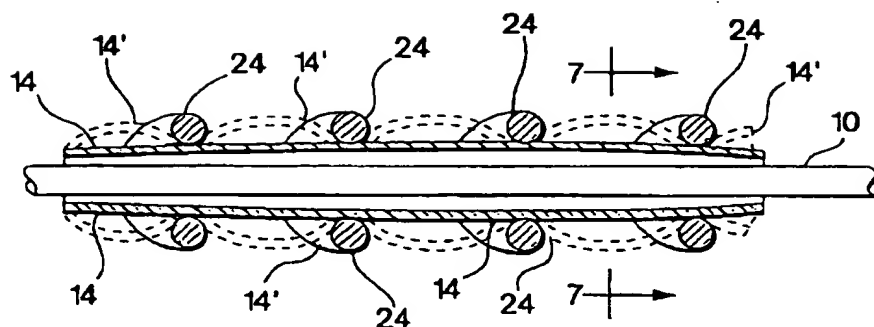


Fig. 5

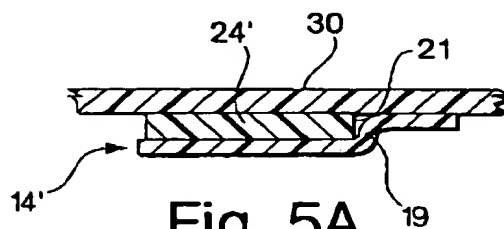


Fig. 5A

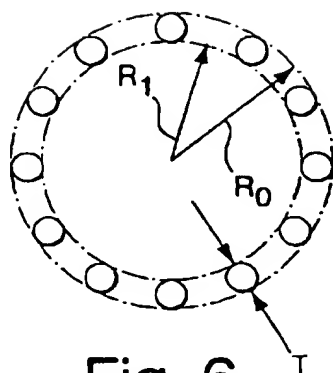


Fig. 6

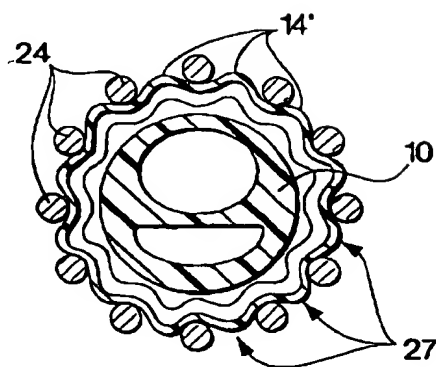


Fig. 7

SUBSTITUTE SHEET (RULE 26)

INTERNATIONAL SEARCH REPORT

In International Application No

PCT/IE 98/00076

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A61F2/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61F A61M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X,P	EP 0 834 293 A (CORDIS EUROPA N.V.) 8 April 1998 see abstract see column 1, line 20 - line 55; claims 1-7; figures 1-4	1-20
X,P	FR 2 753 907 A (LABORATOIRES NYCOMED S.A.) 3 April 1998 see abstract see page 1, line 22 - page 3, line 26; figures 2A,B,3A-E	1-20
X	EP 0 770 366 A (FIGULLA) 2 May 1997 see abstract	1,5-7
A	see column 2, line 20 - line 45; claims 1,4; figures 1-3	2-4,8-20
	--- -/-	



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

1 December 1998

Date of mailing of the international search report

09/12/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Michels, N

INTERNATIONAL SEARCH REPORT

Int ernational Application No

PCT/IE 98/00076

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 778 010 A (APPLIED VASCULAR ENGINEERING. INC.) 11 June 1997	1,5-7
A	see abstract; figures 2-7 ----	2-4,8-20
A	US 5 250 070 A (PARODI) 5 October 1993 see abstract; claims 1-4; figures 1-4,6 -----	1-20

INTERNATIONAL SEARCH REPORT

Information on patent family members

Int l Application No

PCT/IE 98/00076

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0834293 A	08-04-1998	NL 1004162 C CA 2217085 A	02-04-1998 01-04-1998
FR 2753907 A	03-04-1998	WO 9814233 A	09-04-1998
EP 0770366 A	02-05-1997	DE 19540084 A AT 157237 T DE 59600020 D ES 2106645 T	30-04-1997 15-09-1997 02-10-1997 01-11-1997
EP 0778010 A	11-06-1997	AU 7196296 A CA 2190795 A JP 9266950 A	12-06-1997 08-06-1997 14-10-1997
US 5250070 A	05-10-1993	NONE	

Form PCT/ISA/210 (patent family annex) (July 1992)